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Amendments to the Claims:

The listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims

Claims 1-28 (canceled):

Claim 29 (currently amended): A peptide having the structure:

$$Z^{1}$$
- X^{6} - X^{7} - X^{8} - X^{9} - X^{10} - X^{11} - X^{12} - X^{13} - X^{14} - X^{15} - X^{16} - X^{17} - Z^{2}

wherein X⁶ is selected from the group consisting of: D-arginine, D- alanine, D-norleucine, D-α-aminobutyric acid, D-valine, D-leucine, D-isoleucine, D- proline, D-methionine, D- phenylalanine, D- asparagine, D-glutamine, D- serine, D-threonine, D- glutamic acid, D-aspartic acid, D- lysine, D-histidine, D-tryptophan, D-tyrosine, D-cyclohexylalanine, D-(2')naphthylalanine, D-ornithine, D-homoarginine, D-norarginine, D-norarginine, D-citrulline and 5-guanidinopropionic acid,

X⁷ is cysteine,

X8 is either methionine, norleucine, or N-methyl norleucine,

X⁹ is leucine.

X¹⁰ is either asparagine, glutamine, leucine, isoleucine, valine, norleucine, cyclohexylalanine, phenylalanine, (2')-naphthylalanine, tyrosine, histidine, tryptophan, lysine, serine, threonine, methionine, or citrulline,

X11 is arginine,

X12 is valine,

X¹³ is phenylalanine, (2')napthylalanine, p-fluoro-phenylalanine, tyrosine, or cyclohexylalanine,

X¹⁴ is arginine or alanine,

X¹⁵ is either proline or sarcosine,

X16 is cysteine or D-cysteine,

X¹⁷ is an optionally present amino acid that, if present, is either tryptophan or tyrosine,

 Z^1 is an optionally present protecting group that, if present, is covalently joined to the N-terminal amino group,

Z² is an optionally present protecting group that, if present, is covalently joined to the C-terminal carboxy group,

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wherein a substituent said peptide optionally contains on said peptide is optionally substituted with a detectable label,

or a pharmaceutically acceptable salt of said peptide.

Claim 30 (previously presented): The peptide of claim 29, wherein said detectable label is selected from the group consisting of: a luminescent label, an enzymatic label, and a radiolabel.

Claim 31 (previously presented): The peptide of claim 30, wherein said detectable label is not present.

Claim 32 (previously presented): The peptide of claim 30, wherein X⁶ is either D-arginine, D-alanine, D-norleucine, D-proline, D-phenylalanine, D-asparagine, D-serine, D-glutamic acid, D-lysine, or D-citrulline.

Claim 33 (currently amended): The peptide of claim 32, wherein X¹⁰ is glutamine or arginine.

Claim 34 (previously presented): The peptide of claim 33, wherein said peptide is substituted with a radiolabel.

Claim 35 (previously presented): The peptide of claim 33, wherein said peptide is not substituted with a detectable label.

Claim 36 (previously presented): The peptide of claim 31, wherein X^{17} is not present, Z^1 is -C(O)CH₃ and Z^2 is -NH₂.

Claim 37 (previously presented): The peptide of claim 33, wherein X^{17} is not present, Z^1 is -C(O)CH3 and Z^2 is -NH2.

Claim 38 (previously presented): The peptide of claim 29, wherein said peptide consists of a sequence selected from the group consisting of: SEQ ID NOs: 29, 30, 31, 32, 33, and 34.

Claim 39 (previously presented): The peptide of claim 29, wherein said peptide consists of SEQ ID NO: 30.

Claim 40 (currently amended): A method of screening for a compound able to bind MCH-1R comprising the step of measuring the ability of said compound to inhibit binding of a <u>detectably</u> labeled peptide of claim 29 to MCH-1R by measuring the change in <u>detectable label</u>.

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Claim 41 (previously presented): The method of claim 40, wherein said peptide is radiolabeled.